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European Patent Office
Office européen des brevets



(11) Publication number : **0 586 094 A1**

(12)

EUROPEAN PATENT APPLICATION

(21) Application number : **93306068.3**

(51) Int. Cl.⁵ : **C12N 15/12, C12P 21/00,
C07K 15/00**

(22) Date of filing : **30.07.93**

The applicant has subsequently filed a sequence listing and declared, that it includes no new matter.

(30) Priority : **03.08.92 US 925403**

(43) Date of publication of application :
09.03.94 Bulletin 94/10

(84) Designated Contracting States :
**AT BE CH DE DK ES FR GB GR IE IT LI LU MC
NL PT SE**

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(54) **Soluble LDL receptor and gene.**

(57) A truncation of the human LDL receptor gene has been created which can be expressed in a heterologous host to produce a soluble fragment to the receptor protein. The gene can be expressed in insect cells in culture to produce a protein fragment which is not only water soluble but which also retains affinity for binding to LDL. The truncation is a truncation of the carboxyl terminus of the native LDL receptor gene which results in a 354 amino acid protein fragment designated LDL-R-³⁵⁴.

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Field of the Invention

The present invention relates to the general field of molecular biology and relates, in particular, to the isolation and utilization of novel gene fragments to create useful fragments of proteins for therapeutic and research purposes.

Background of the Invention

Cholesterol is an abundant substance in animals because it has an essential function in the membranes of animal cells. Cholesterol is almost completely insoluble in aqueous solution. Within the body other than in membranes, therefore, cholesterol must be carefully segregated from exposure to aqueous environments. Therefore, in multicellular organisms, cholesterol is transported about the body packaged within the hydrophobic cores of plasma lipoproteins. The various plasma lipoproteins in humans have been classified into four major classes, principally by buoyant density. The four major classes are very low density lipoprotein (VLDL), intermediate density lipoprotein (IDL), low density lipoprotein (LDL), and high density lipoprotein (HDL). A fifth class of lipoprotein is chylomicrons, which occur only after feeding. The most abundant cholesterol-carrying lipoproteins in human plasma are of the LDL class, and high levels of LDL circulating in blood plasma are associated with arteriosclerosis and coronary dysfunction. Hence, the mechanisms which purge LDL from the blood stream have been subject to significant study.

In higher animals, LDL is scavenged from the blood stream by receptors located on the cell membranes of animal cells. The receptors are known as the LDL receptor. The structure of the LDL receptor has been generally described, and the gene coding for the production of the receptor has been isolated and sequenced. Descriptions of the LDL receptor, its structure, and its general functionality in interaction with other cellular processes can be found in Schneider, *Bio. Et. Biophys. Acta.*, 988, pages 307-317 (1989) and Hobbs, et al., *Annu. Rev. Genet.*, 24, pages 133-170 (1990). The LDL receptor protein is a cell surface glycoprotein that regulates plasma cholesterol by mediating endocytosis of lipoproteins. The human LDL receptor is a protein of 860 amino acids encoded by a gene which actuates the transcription of an mRNA of 5.3 kb in length. The mRNA of the human LDL receptor gene includes a long 3' untranslated region, as well as a signal peptide which actuates transport of the protein to the plasma membrane, and which is cleaved from the mature form of the protein.

The LDL receptor protein has been characterized as having generally five different domains or regions. One region, referred to as the ligand binding domain, is generally associated with the affinity of the receptor for the two known proteins which are bound by the receptor, ApoB-100, the 550 kd glycoprotein that is the predominant protein of LDL, and ApoE, a 34 kd protein found in multiple copies in IDL and a subclass of HDL. Other domains in the mature LDL receptor protein include an EGF (epidermal growth factor) precursor homology region which, while not specifically necessary for ligand affinity, appears to play a role in the presentation of the LDL receptor to lipoproteins external to the cell. The third domain of the LDL receptor is a region containing clustered O-linked sugar chains. A membrane spanning region of 22 amino acids, of high hydrophobicity, is responsible for the membrane bound nature of the protein. At the carboxyl end of the protein is a domain of 50 amino acids, which extends into the cytoplasm.

While the human LDL receptor gene and the human LDL receptor protein have been studied extensively, this study has been a difficult and, at times, an arduous process. One of the reasons for the difficulty of this study is the inability of researchers to isolate and utilize large quantities of the LDL receptor protein itself. While the protein is produced in significant quantities in animal cells, it is membrane bound and presumably insoluble in aqueous solution. Hence, it is quite difficult to isolate the receptor protein from mammalian tissues in any significant quantity. In addition, since the affinity for membranes is quite strong, it is difficult to express the human LDL receptor protein in heterologous hosts and isolate pure quantities of protein, due to the strong tendency of the protein to associate with membranes in the expression system. In addition, the protein has a complex three dimensional structure, particularly in the ligand binding domain, which is cysteine-rich and includes multiple disulfide bonds. Since the three dimensional structure of the protein is essential for the affinity of the receptor to LDL, production of the protein in hosts which do not properly process proteins subsequent to translation to result in a correct three dimensional structure would not give rise to functional receptor protein.

Summary of the Invention

The present invention is summarized in that a truncated portion of the human LDL receptor gene has been identified which can be expressed in insect cells in culture to produce a truncated water-soluble form of the LDL receptor protein which binds to, and has specific affinity for, LDL to which the native receptor has affinity.

It is an object of the present invention to describe a gene fragment, and the protein produced from this fragment, which enables the production and accumulation of significant quantities of an LDL receptor fragment which has affinity for and recognizes LDL, and yet which is soluble in aqueous solutions.

It is another object of the present invention to provide a new substance of possible therapeutic value, specifically a soluble LDL receptor protein fragment which can be produced in heterologous hosts for possible clinical, diagnostic, and research use on human systems.

It is a further object of the present invention to elucidate and describe a mechanism by which the active binding portion of the LDL receptor can be created in heterologous hosts, so as to be useful for further research purposes in further investigating the interaction of the receptor and its target molecules.

It is yet another object of the present invention to describe possible practical applications made possible by the expression of the water-soluble LDL receptor fragment described here.

Other objects, advantages, and features of the present invention will become apparent from the following specification when taken in conjunction with the accompanying drawings.

Brief Description of the Drawings

Figure 1 is a schematic view of the current model for a human LDL receptor protein.

Figure 2 is a model of the predicted schematic structure of the truncated LDL receptor LDL-R³⁵⁴.

Figure 3 is a graphical representation of a competition binding assay verifying the binding of the LDL receptor produced in accordance with the present invention with LDL.

Detailed Description of the Invention

In accordance with the present invention, it has been found that a truncated portion of the gene for human low density lipoprotein receptor (LDL-R) enables the production of a soluble form of the LDL receptor which retains the binding specificity of the native LDL receptor. The expression of this truncated gene in insect cells in culture results in the production of a truncated LDL receptor protein which appears to be conformationally correct, and which, although soluble in aqueous solution, exhibits proper specificity for LDL proteins so as to be useful for LDL binding.

The mechanism of the present invention may be best understood by referring first to Figure 1 which is a graphical illustration of the currently best understood model of the LDL receptor, and the various domains which constitute that protein. The functional domains of the LDL receptor protein had been identified by virtue of homology to other proteins, and also by virtue of the phenotypes of naturally occurring mutants of the protein found in the human population, which have been extensively studied by Brown and Goldstein. Five domains of the mature protein have been characterized. The first is an N-terminal cysteine-rich domain, which is associated with ligand binding, and referred to as the ligand binding domain in Figure 1. The next region in the protein is a region having homology to the epidermal growth factor (EGF) precursor. The EGF precursor homology domain is required for the acid-dependent disassociation of the lipoproteins from the receptor, during receptor recycling, and also serves to position the ligand-binding domain so that it can bind LDL on the cell surface. The third domain of the protein is a region rich in O-linked carbohydrate. The next region of the protein is a membrane spanning domain, of extremely hydrophobic character, which is the portion of the protein responsible for binding to the animal cell membrane. A cytoplasmic domain at the extreme carboxyl terminus of the protein completes the regions of the protein.

It is previously known that virtually all polymorphisms in the human receptor that are associated with defective binding of LDL reside in the extreme N-terminal cysteine-rich domain. Based on the mutagenic analysis of the LDL receptor as performed by Esser, et al. (*J. Biol. Chem.*, 263:26, pages 13282-13290 (1988)) and Russell, et al. (*J. Biol. Chem.*, 264:36, pages 21682-21688 (1989)), it is clear that all of the seven imperfect repeats which characterize the ligand binding domain are required, or at least involved, in optimal binding to the LDL ligand. Two additional imperfect repeats, designated A and B in Figure 1, are also found in the EGF precursor homology domain. It seems likely from a similar mutation analysis that repeat A may also facilitate optimal ligand binding.

Based on this understanding, it was decided to attempt a truncation of the full-length LDL receptor gene, to isolate a protein coding sequence encoding for the amino terminal portion of the protein, which would include the expression of all repeats 1 through 7 of the ligand binding domain plus repeat A of the region of EGF precursor homology region. If a host cell expressing this truncated gene were able to facilitate the proper folding of the truncated protein product, then it was possible that the truncated product would retain its ability to bind LDL. Furthermore, the exclusion of the membrane spanning domain could possibly result in secretion of the produced protein. Prior to this attempt, it was not known if a conformationally correct ligand binding domain

could be produced without the rest of the native protein. Formation of the correct structure of complex proteins is still poorly understood.

It was previously known that a naturally-occurring mutation of the LDL receptor that deletes the membrane spanning and cytoplasmic domains results in the partial secretion of a truncated receptor, rather than the localization of all of the truncated receptor onto the membrane. However, it is not known if any biological activity of this extracellular mutant receptor has been detected.

By utilization of a restriction endonuclease cleavage site in the commonly available LDL receptor cDNA clone, it was possible to prepare a gene fragment encoding the first 354 amino acids of the mature form of the human LDL-R protein. The polypeptide thus created, designated here LDL-R³⁵⁴, could be produced in a foreign host and retain specificity for the LDL molecule. The LDL-R³⁵⁴ is also soluble in aqueous solution, and therefore has independent utility in studies of binding to LDL.

As described below, the truncated LDL receptor gene fragment coding for LDL-R³⁵⁴ has been introduced, through the use of a recombinant baculovirus, into insect cells grown in culture. It is believed that the utilization of eukaryotic cells in general, and insect cells in particular, is an important factor in the successful production of significant quantities of receptor binding soluble protein. Because the conformational structure of the receptor is critical to its ability to bind to the LDL proteins, if the polypeptide could not be produced in a manner that yielded a properly folded product in heterologous host cells, effective binding would not be possible. It is likely that other eukaryotic host cell systems will result in production of the same protein fragment. However, since the production of proteins of correct conformational character cannot be predicted, particularly for truncated forms of protein, it was not known in advance if a conformationally correct receptor, i.e. one having affinity to the target molecules, could be produced in any host. Nevertheless, the structure of the protein fragment expressed here does seem correct. There is also reason to believe that this particular truncation, with the possible exception of an amino terminal deletion of the very first repeat sequence, is probably the most minimal truncation that would produce active protein. Other truncations which are less severe are likely, based on the evidence presented here, to also express conformationally correct protein fragments. Any such larger truncations could include the EGF precursor homology region without affecting ligand binding, but should not include the hydrophobic membrane spanning domain from the native gene.

The cDNA for human LDL receptor has previously been isolated by others, sequenced, and published. A copy of the 5.3 kb cDNA has been deposited by others with the ATCC and hosted in *E. coli* at Accession Number 57004. From that vector, the region encoding the critical amino terminal 354 amino acid polypeptide can readily be isolated, by a procedure similar to that described below.

Applications of Truncated LDL Receptor

One category of use of the truncated LDL receptor protein is for diagnosis. Presently used diagnostic assays of human serum LDL levels are performed by an indirect estimate method based on a precipitating polyanion used to precipitate LDL with VLDL. The amount of cholesterol in VLDL is estimated by dividing total plasma triglyceride by 5. This method only estimates LDL cholesterol and is incapable of identifying patients who have greater numbers of LDL particles, but only modest hypercholesterolemia, because their LDL particles are simply smaller than the norm, i.e. with less cholesterol per particle. These patients may be even more at risk if, as some data suggest, smaller LDL particles are actually more atherogenic than larger particles. Thus a method of measuring either or both of protein and lipid content of the LDL would be highly desirable. Information as to both lipid and protein would enable estimates to be made of the mean size of LDL particles, which appears to be a clinically useful indication. The truncated LDL receptor described here could be used in binding assays against blood or serum to ascertain actual protein levels and/or numbers of LDL particles.

Another potential advantage of a method of specifically capturing lipoprotein particles containing apoB from the blood would be potential quantitation of Lipoprotein (a), or Lp(a). Lp(a) is a protein covalently attached to the major LDL protein (apoB), and the plasma level of Lp(a) has proven to have a strong correlation with heart disease risk. The plasma level of Lp(a) appears to be both extremely variable and completely determined by genetic factors, and the level of Lp(a) results in a protein product that can vary in size by as much as 20-fold. The immunoassays currently used to quantitate serum Lp(a) are not valid for absolute (i.e. size independent) quantitation. The development of a protocol for binding LDL through the use of the LDL receptor described here, may permit Lp(a) to be measured, since Lp(a) is easily released from apoB by reducing agents which break the linking disulfide bonds.

Another application of the soluble LDL receptor would be for the isolation of plasma LDL for research or clinical purposes. Presently LDL is isolated by preparative ultracentrifugation, which is very costly and requires six days of continuous ultracentrifugation time. A commercially available method or kit for LDL isolation based on this truncated receptor would be popular among LDL researchers, particularly since the receptor recognizes

apoB only on LDL and not on VLDL. An advantageous feature of the use of the truncated LDL receptor for affinity selection, rather than an antibody to apoB, is both this selectivity for LDL and also the fact that although the LDL-LDL receptor affinity is quite high, the affinity depends on calcium and can readily be dissipated through the use of a chelating agent such as EDTA. Antibodies to apoB typically require much harsher elution conditions.

Another more dramatic possible application for this receptor would be essentially blood filtering for LDL. The immobilized truncated LDL receptor produced in quantity can be immobilized in a column or cartridge through which a patient's blood could be passed to specifically remove LDL from the blood. The column or cartridge could be regenerated by eluting the LDL. This therapeutic approach seems particularly adapted to kidney dialysis patients, who invariably suffer from hyperlipidemia and who already have their blood dialyzed on a regular basis.

It is also possible to construct alternate LDL-R³⁵⁴ expression vectors which condition secretion of the expressed protein fragment. Since the mature protein is initially produced with a signal peptide, which causes transportation of the immature form of the protein to the plasma membrane of the cell which produces the protein, there is every reason to expect that a similar signal peptide strategy can be used to cause secretion of the truncated form of the protein described herein.

EXAMPLES

Construction of Gene.

The construction of the coding sequence for the LDL-R³⁵⁴ polypeptide began with a vector designated pRGLDLR-13, which consists of a 1317 base pair XbaI, RsaI fragment from the plasmid pDLR3 which has been deposited by others with the ATCC as Accession No. 57004. The vector pRGLDLR-13 thus includes a sequence encoding the first 425 amino acids of the LDL receptor, which has been inserted by blunt end ligation into the EcoRV site of pBluescript M13+ (Stratagene) 3'-5' orientation. A 14 base pair oligonucleotide (CTAGTCTAGACTAG, New England Biolabs) was used to introduce a translational termination codon in all reading frames, by insertion at the unique EcoRV site in pRGLDLR-13. The EcoRV site corresponds to nucleotide 1076 of the LDL-R cDNA. The resulting plasmid was designated pDL-LDLR. The 5' and 3' termini of the approximately 1317-bp LDL-R fragment from pDL-LDLR was confirmed by DNA sequence analysis. The translation terminating codon was in the predicted reading frame. An 1100 pb BamHI fragment of pDL-LDLR was then ligated to copies of a plasmid pAcYM1, which had been digested with BamHI and treated with alkaline phosphatase. The details of pAcYM1 are described in Matsuura et al. *J. Gen. Virol.* 68:1233-1250 (1987). The resulting construction was designated pAcLDL-R³⁵⁴. This transfer vector thus consists of the 5' and 3' sequences from the *Autographa californica* polyhedrin gene which are flanking a protein coding sequence of nucleotides which encodes the first 354 residues of the human LDL receptor, followed by two additional amino acids (leucine and valine), which were introduced at the carboxyl terminus of the peptide by oligonucleotide insertion.

Production of Recombinant Baculovirus.

Copies of the plasmid pAcLDL-R³⁵⁴ constructed as described above were co-transfected with the wild-type baculovirus genome (AcNPV) into *Spodoptera frugiperda* Sf-21 cells. The recombinant viruses (AcNPV-R³⁵⁴) harboring the sequence for the truncated LDL receptor were selected, plaque purified, and titered as described in Summers et al., *A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures*, Tex. Agric. Exp. Stn. Bull. (1987) and Miller et al. in *Genetic Engineering*, Plenum, New York, pp. 277-298 (1986).

Recombinant virus (AcNPV-R³⁵⁴) served as the template for a polymerase chain reaction utilizing amplifying oligonucleotides homologous to the 5' and 3' sequences of the *Autographa californica* polyhedrin gene. The resulting amplification product was of a size consistent with the predicted recombination event.

Transfection of Cultured Insect Cells and Recovery of Protein.

The recombinant baculoviruses were then used to infect cells of SF-21 in culture. Quantities of the recombinant baculovirus were infected onto dishes of 4×10^6 SF-21 cells at a multiplicity of infection of four. Two milliliters of SF-900 (Gibco) medium were harvested from the infected cells 52 hours after infection with the recombinant baculovirus in TCA precipitated media to yield 60 μ gram of protein. Media samples were subjected to sodium dodecyl sulfate, polyacrylamide gel electrophoresis (SDS-PAGE), in the absence or presence of beta-mercaptoethanol, with non-reducing gels of 13.3% acrylamide resolving gel with a 6% stacker and with

reducing gels of 10% acrylamide resolving gel with a 4.5% stacker. Immunoblotting was performed using anti-human LDR-R monoclonal antibody C7, and alkaline phosphatase-conjugated goat anti-mouse IgG (Sigma).

Binding Experiments.

Media from transfected SF-21 cells was collected 52 hours after infection and was dialyzed against PBS at pH 7.4. Separately, CHO cells [line TR715-19] previously demonstrated to overexpress the human LDL receptor were grown in Hams nutrient mixture F-12 (Gibco), 20 mM Hepes, pH 7.2, 1% (V/V) fetal calf serum, 4% (V/V) lipoprotein deficient serum (LDS), 2 mM glutamine, and 100 units per milliliter penicillin, 100 µg per ml streptomycin. Twelve hours prior to use, confluent cells were switched to a medium containing Hams F-12, 20 mM Hepes pH 7.2, 5% LDS, 2 mM glutamine, 100 units per ml penicillin, 100 µg per ml streptomycin, 10 µg per ml cholesterol and 0.1 µg per ml 25-hydroxycholesterol. The indicated concentration of dialyzed tissue culture proteins (the concentration was determined by a modified Lowry assay) were incubated with 1.0 µg of [¹²⁵I]-LDL in 2 ml Hams F-12, 20 mM Hepes, pH 7.2 and 5 mg/ml bovine serum albumin, at ambient temperatures for 90 minutes. Samples were equilibrated to 4°C prior to addition to the CHO cells. The cells were incubated for 3 hours at 4°C on a rotary shaker, prior to washing, and the release of bound [¹²⁵I]-LDL by dextran sulfate. Radio iodination of LDL was performed as described previously, e.g. Checovich et al., *Biochem.*, 27:1934 (1988).

The results as illustrated in Fig. 3 indicate that for increasing concentrations of culture media incubated with radiolabeled LDL, in the presence of cells expressing the LDL receptor, the ability of the LDL-R³⁵⁴ to specifically bind to LDL was demonstrated. The decrease in cell associated radiolabeled LDL in the presence of increasing concentrations of proteins from the medium of truncated receptor expressing cells demonstrated competitive LDL binding by the protein fragments in the medium. Competitions of similarity severity were not observed with equivalent concentrations of proteins from wild-type virus infected cells.

These results of the immunoblots performed as described above indicate that the peptide fragment which is produced in the infected insect cells described above is secreted, since it is contained with the media extracted from the culture and recognized by the LDL-R specific monoclonal antibody C7. Furthermore, the competitive binding assay indicates that the fragment does correctly bind to the LDL molecule, with a specificity that is at least competitive with the native membrane-bound LDL receptor that is produced in CHO cells. Thus, for the first time, a soluble form of the LDL receptor has been created, and is now made available for use in non-membrane bound research, diagnostic, and therapeutic applications.

5

SEQUENCE LISTING

10

(1) GENERAL INFORMATION:

(i) APPLICANT:

NAME: Wisconsin Alumni Research Foundation
 STREET: P O Box 7365
 CITY: Madison, Wisconsin 53707
 COUNTRY: U S A

15

(ii) TITLE OF INVENTION: Soluble LDL Receptor and Gene

(iii) NUMBER OF SEQUENCES: 2

20

(iv) COMPUTER READABLE FORM:

- (A) MEDIUM TYPE: Floppy disk
- (B) COMPUTER: IBM PC compatible
- (C) OPERATING SYSTEM: PC-DOS/MS-DOS
- (D) SOFTWARE: PatentIn Release #1.0, Version #1.25 EPO

25

(v) CURRENT APPLICATION DATA:

APPLICATION NUMBER: EP 93306068.3

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1317 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

30

(ii) MOLECULE TYPE: cDNA to mRNA

(iii) HYPOTHETICAL: NO

35

(iv) ANTI-SENSE: NO

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

40

(vii) IMMEDIATE SOURCE:

(B) CLONE: LDL Receptor

(ix) FEATURE:

- (A) NAME/KEY: CDC
- (B) LOCATION: 16..1086

45

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

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1

5

10

50

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55

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	GGC CGT GTC AAC CGC TGC ATT CCT CAG TTC TGG AGG TGC GAT GGC CAA Gly Arg Val Asn Arg Cys Ile Pro Gln Phe Trp Arg Cys Asp Gly Gln 80 85 90	291		
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25	TGC TCC CAG GAC GAG TTT CGC TGC CAC GAT GGG AAG TGC ATC TCT CGG Cys Ser Gln Asp Glu Phe Arg Cys His Asp Gly Lys Cys Ile Ser Arg 110 115 120	387		
	CAG TTC GTC TGT GAC TCA GAC CGG GAC TGC TTG GAC GGC TCA GAC GAG Gln Phe Val Cys Asp Ser Asp Arg Asp Cys Leu Asp Gly Ser Asp Glu 125 130 135 140	435		
30	GCC TCC TGC CCG GTG CTC ACC TGT GGT CCC GCC AGC TTC CAG TGC AAC Ala Ser Cys Pro Val Leu Thr Cys Gly Pro Ala Ser Phe Gln Cys Asn 145 150 155	483		
	AGC TCC ACC TGC ATC CCC CAG CTG TGG GCC TGC GAC AAC GAC CCC GAC Ser Ser Thr Cys Ile Pro Gln Leu Trp Ala Cys Asp Asn Asp Pro Asp 160 165 170	531		
35	TGC GAA GAT GGC TCG GAT GAG TGG CCG CAG CGC TGT AGG GGT CTT TAC Cys Glu Asp Gly Ser Asp Glu Trp Pro Gln Arg Cys Arg Gly Leu Tyr 175 180 185	579		
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55

5

CGG CAG TGT GAC CGG GAA TAT GAC TGC AAG GAC ATG AGC GAT GAA GTT 819
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 270 275 280

CAC AGC GGC GAA TGC ATC ACC CTG GAC AAA GTC TGC AAC ATG GCT AGA 915
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 285 290 295 300

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 305 310 315

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GCCGGGGGTC CGCTAGTTCT AGAA 1317

35 (2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 356 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

40

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

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 20 25 30

Gln Cys Gln Asp Gly Lys Cys Ile Ser Tyr Lys Trp Val Cys Asp Gly
 35 40 45

50 Ser Ala Glu Cys Gln Asp Gly Ser Asp Glu Ser Gln Glu Thr Cys Leu

55

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		130	135 140
20	Val Leu Thr Cys Gly Pro Ala Ser Phe Gln Cys Asn Ser Ser Thr Cys		
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	Ile Pro Gln Leu Trp Ala Cys Asp Asn Asp Pro Asp Cys Glu Asp Gly		
		165	170 175
25	Ser Asp Glu Trp Pro Gln Arg Cys Arg Gly Leu Tyr Val Phe Gln Gly		
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30	Cys Ile His Ser Ser Trp Arg Cys Asp Gly Gly Pro Asp Cys Lys Asp		
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	Lys Ser Asp Glu Glu Asn Cys Ala Val Ala Thr Cys Arg Pro Asp Glu		
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	Arg Glu Tyr Asp Cys Lys Asp Met Ser Asp Glu Val Gly Cys Val Asn		
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	Asn Asn Gly Gly Cys Ser His Val Cys Asn Asp Leu Lys Ile Gly Tyr		
		325	330 335
50	Glu Cys Leu Cys Pro Asp Gly Phe Gln Leu Val Ala Gln Arg Arg Cys		
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	Glu Asp Leu Val		
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55

(I. INFORMATION FOR SEQ ID NO:3:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 14 base pairs
 (B) TYPE: nucleic acid
 (C) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA probe

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

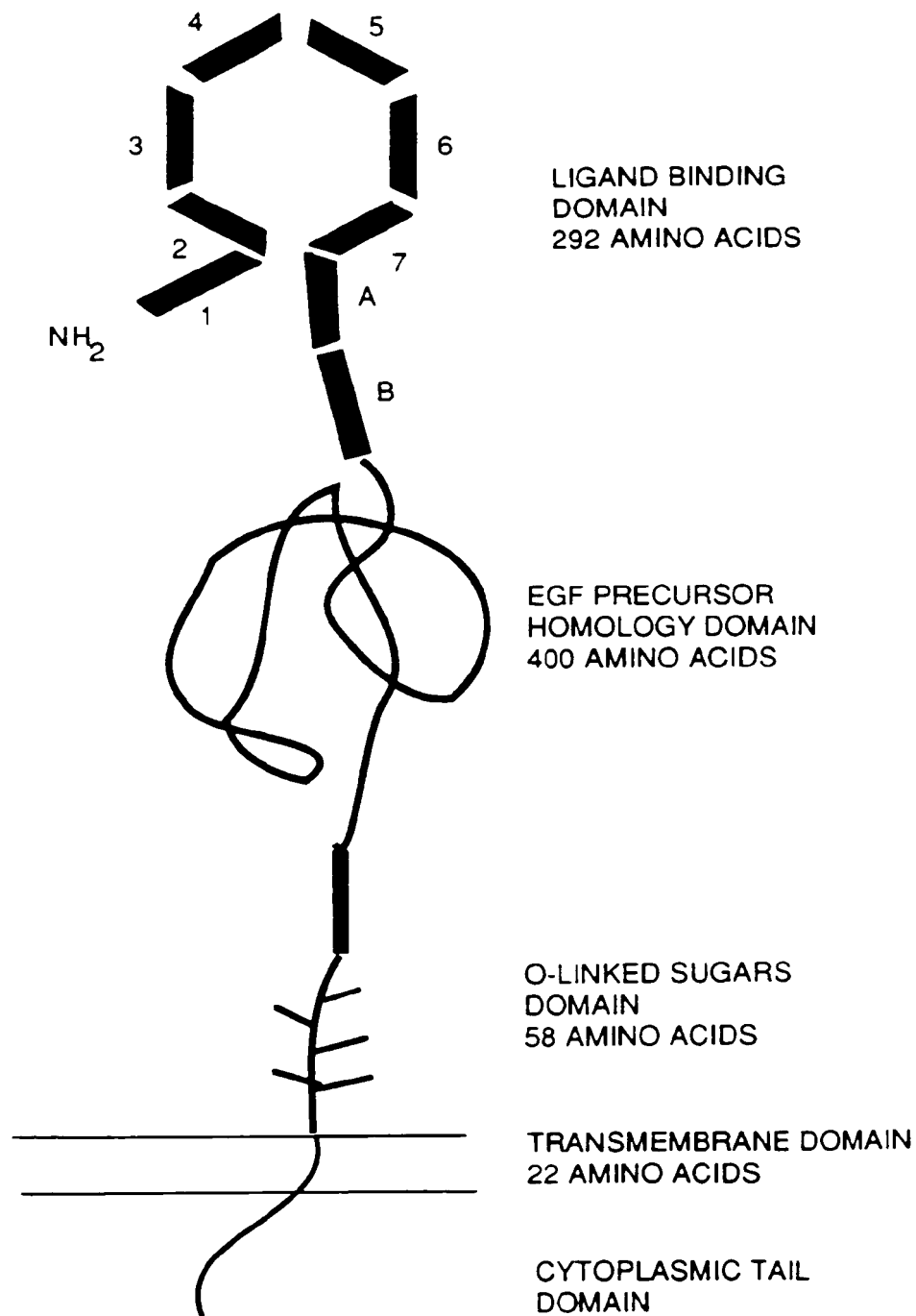
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Claims

1. A water soluble polypeptide having LDL receptor binding affinity for low density lipoprotein (LDL).
2. A polypeptide according to claim 1 comprising a truncated LDL receptor protein.
3. A polypeptide according to claim 2 comprising a ligand binding domain and a portion of an epidermal growth factor homology domain of the LDL receptor protein.
4. A polypeptide according to claim 3 comprising a truncation after amino acid 354 of the LDL receptor protein.
5. A polypeptide according to claim 3 or 4 which consists essentially of a truncation of native LDL receptor protein.
6. A polypeptide according to claim 4 or 5 wherein the truncated LDL receptor polypeptide consists of amino acid residues 1 to 354 of LDL receptor protein.
7. A method of making a water soluble polypeptide according to any one of claims 1 to 6, in a host comprising the steps of
 - creating a nucleotide sequence comprising a coding sequence coding for a truncated form of a LDL receptor gene which does not include the membrane spanning domain, and which further comprises flanking regulatory sequences effective to express the coding sequence in cells of the host;
 - transforming the nucleotide sequence into cells of the host;
 - culturing the cells in a medium under conditions effective to produce the polypeptide; and recovering the polypeptide from the cultured cells.
8. A method according to claim 7 in which the host is heterologous and eukaryotic and host cells are cultured under conditions effective to secrete the truncated LDL receptor polypeptide.
9. A method according to claim 8 wherein the heterologous host is insect cells in culture.
10. A method according to claim 10 wherein the insect cells are Sf-21 cells.
11. A method according to any one of claims 7 to 10 wherein the nucleotide sequence is introduced into the host cells through the use of a baculovirus.

FIG. 1



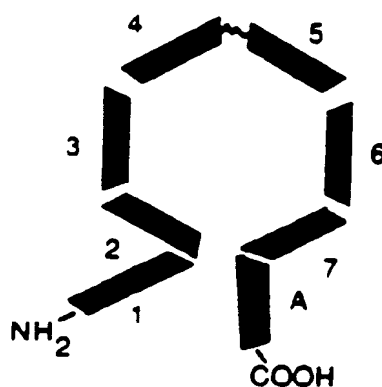


FIG. 2

Competition for LDL Binding:
WT vs LDL receptor media

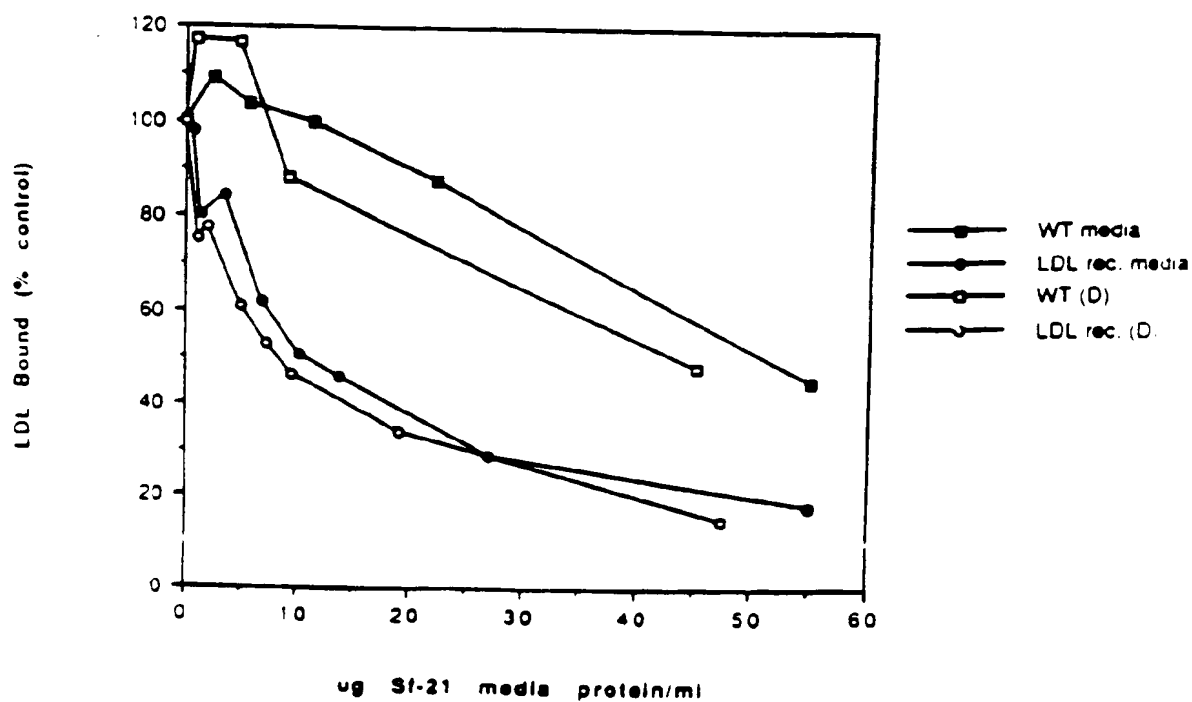


FIG. 3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

DOCUMENTS CONSIDERED TO BE RELEVANT			EP 93306068.3
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
X	WO - A - 86/04 090 (BOARD OF REGENTS, THE UNIVERSITY OF TEXAS SYSTEM) * Claims 16-21; fig. 3B *	1	C 12 N 15/12 C 12 P 21/00 C 07 K 15/00
X, D	THE JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 263, no. 25, September 05, 1988, Baltimore, USA V. ESSER et al. "Mutational Analysis of the Ligand Binding Domain of the Low Density Lipoprotein Receptor" pages 13282-13290 * Totality *	1	
X, D	THE JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 264, no. 35, December 15, 1989, Baltimore, USA D.W. RUSSELL et al. "Different Combinations of Cysteine-rich Repeats Mediate Binding of Low Density Lipo- protein Receptor to Two Different Proteins" pages 21682-21688 * Totality *	1	TECHNICAL FIELDS SEARCHED (Int. Cl.5) C 12 N C 12 P C 07 K
A	WO - A - 90/05 748 (MASSACHUSETTS INSTITUTE OF TECHNOLOGY) * Abstract *	1	
A	WC - A - 87/02 061 (BIO-TECHNOLOGY RESEARCH PARTNERS) * Claims *	1	
D, A	BIOCHIMICA ET BIOPHYSICA		
The present search report has been drawn up for all claims			
Place of search VIENNA		Date of completion of the search 05-11-1993	Examiner WOLF
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

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Application Number

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EP 93306068.3

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